



#9
PATENT *12*
1/14/03

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of: ANNAPRAGADA et al.

Attorney Docket No.:
LAM1P154/P0696

Application No.: 09/688,021

Examiner: ANDERSON, Matthew A.

Filed: October 13, 2000

Group: 1765

Title: PROCESS FOR ETCHING VIAS IN
ORGANOSILICATE GLASS MATERIALS
WITHOUT CAUSING RIE LAG

RECEIVED
JAN 13 2003
TC 1700

CERTIFICATE OF MAILING

I hereby certify that this correspondence is being deposited with the
United States Postal Service as First Class Mail to: Box AF,
Commissioner for Patents, Washington, D.C. 20231 on January 3, 2003.

Signed: *Sue Funchess*

Sue Funchess

AMENDMENT B

Box AF
Commissioner for Patents
Washington, D.C. 20231

Dear Sir:

In response to the Office communication dated November 20, 2002, please consider the
following:

REMARKS

The Examiner rejected claims 1 and 7 under 35 U.S.C. 102(b) as being anticipated by
Hung et al. (US 6,387,287 B1). The Examiner stated that Hung et al. discloses in col. 16, lines
1-43 and in Table 10 a method of etching with plasma an organic silicate glass (i.e. the TEOS
ARC layer) on a wafer with a gas comprising C₄F₈ and CF₄ and argon, and that in col. 17, lines
20-35, of Hung further suggests the improvement of nitride corner selectivity by the inclusion of
a more polymerizing gas such as CH₂F₂, and that the problems of reduced etch stop can be
counteracted by the use of N₂ or O₂, and that Hung discloses an SiN etch using CH₂F₂, O₂, and
Ar in FIG. 11 and that the need for gas chemistry control points one of ordinary skill to the
inherent placement of a wafer in a reaction chamber for performance of this process.

Claims 1 and 7 do not recite an organic silicate glass as stated by the Examiner, but instead an organosilicate glass. TEOS oxide of Hung is not an organosilicate glass. TEOS oxide is a silicon oxide grown under PECVD using tetraorthosilicate (TEOS) as discussed on col. 2, lines 43-45, of Hung. Since TEOS oxide is a silicon oxide, not organosilicate glass, Hung does not disclose etching a feature in an integrated circuit incorporating at least one layer of organosilicate glass or etching through the organosilicate glass, as recited in claims 1 and 7. For at least these reasons, claims 1 and 7 are not anticipated by Hung.

The Examiner rejected claims 2-5 and 8-10 under 35 U.S.C. 103(a) as being unpatentable over Hung et al. (US 6,387,287 B1). Claims 2-5 and 8-10 are ultimately dependent on claims 1 or 7, and are therefore respectfully submitted to be patentable over the art of record for at least the reasons set forth above with respect to claims 1 and 7. Additionally, these dependent claims require additional elements that, when taken in the context of the claimed invention, further patentably distinguish the art of record. For example, claims 2 and 8 recite that the etchant gas mixture further comprises CH₂F₂. Hung in Table 10 teaches using C₄F₈ and CF₄, but not CO or O₂ for an ARC open, and then C₄F₆, O₂, CO, and Ar, but not C₄F₈ and CF₄, for the main etch. It would not be obvious to add CH₂F₂ to C₄F₈ and CF₄, since the nitride corner selectivity provided by CH₂F₂ is for the main etch, which does not use C₄F₈ and CF₄. The Examiner did not point out why it would be desirable to have nitride corner selectivity during the opening of the ARC, where the nitride corners are not exposed. In addition, the Examiner has not pointed out anything that suggests that CH₂F₂ would successfully provide nitride corner selectivity when opening the ARC, since Hung only teaches that CH₂F₂ provides nitride corner selectivity when performing the main etch. In addition, claims 3 and 9 recite that the etchant gas mixture further comprises oxygen. Again Table 10 of Hung teaches away from using C₄F₈, CF₄ and oxygen in the same etchant gas. Instead, Table 10 of Hung clearly shows that C₄F₈ and CF₄ are stopped when oxygen is flowed. In addition, claim 5 recites that C₄F₈, CF₄, CH₂F₂, oxygen and argon are used to etch the organosilicate glass dielectric. Nothing in Hung suggests that such an etchant gas would be successful in etching organosilicate glass. For at least these reasons, claims 2-5 and 8-10 are not made obvious by the cited references.

The Examiner rejected claims 6 and 11-13 under 35 U.S.C. 103(a) as being unpatentable over Hung as applied to claims 1-5, 7-10 above and further in view of Chiang et al. (US 5,739,579) and Wolf et al. (Volume 1, pp 556). Claims 6 and 11-13 are ultimately dependent on claims 1 or 7, and are therefore respectfully submitted to be patentable over the art of record for

at least the reasons set forth above with respect to claims 1 and 7. Additionally, these dependent claims require additional elements that, when taken in the context of the claimed invention, further patentably distinguish the art of record. For at least these reasons, claims 6 and 11-13 are not made obvious by the cited references.

The Examiner rejected claim 14 under 35 U.S.C. 103(a) as being unpatentable over Hung et al., Chang, and Wolf et al. as applied to claims 1-13 above in view of Li et al. (US 6,284,149 B1). Claim 14 is ultimately dependent on claim 7, and is therefore respectfully submitted to be patentable over the art of record for at least the reasons set forth above with respect to claim 7. Additionally, this dependent claim requires additional elements that, when taken in the context of the claimed invention, further patentably distinguish the art of record. For at least these reasons, claim 14 is not made obvious by the cited references.

In view of the above, it is respectfully submitted that the application is in a condition for allowance and action to that effect is respectfully requested at an early date. If the Examiner feels that a telephone conference would expedite allowance of this application, the Examiner is invited to call the undersigned at (831) 655-2300.

The Commissioner is authorized to charge any fees that may be due to our Deposit Account No. 50-0388 (Order No. LAM1P154).

Respectfully submitted,

BEYER WEAVER & THOMAS, LLP



Michael Lee
Reg. No. 31,846

P.O. Box 778
Berkeley, CA 94704-0778
(831) 655-2300